

In the Claims

1. (currently amended) A composition for transdermal administration of at least one therapeutically active compounds and/or nutrients, which comprises compound or nutrient, said composition comprising:
  - [[(a)]] at least one item selected from the group consisting of
    - at least one therapeutically active compound and and/or at least one nutrient[[,]]; and
  - [[(b)]] a non-oily emulsion.
2. (currently amended) The composition Composition for transdermal administration according to claim 1, characterised in that the wherein said at least one therapeutically active compound and [[or]] said at least one nutrient is an ionic compound.
3. (currently amended) The composition Composition for transdermal administration according to claim 2, characterized in that wherein the ionic compound is a metal ion.
4. (currently amended) The composition Composition according to claim 1, characterised in that the wherein said at least one therapeutically active compound is a polypeptide.
5. (currently amended) The composition Composition according to claim 4, characterised in that the wherein said polypeptide has a molecular weight of up to 7000 kDa.
6. (currently amended) The composition Composition according to claim 1, characterised in that the wherein said at least one therapeutically active compound is selected from the group consisting of [[an]] antiparasitic agents agent, anthelmintic drugs [[or]] and antibiotic drugs [[drug]], used for the treatment of humans, livestock or domestic animals.
7. (currently amended) The composition Composition according to any one of the proceeding claims, characterised in that the claim 1, wherein said non-oily emulsion is a mixture of lecithin, bile salt and cholesterol.
8. (currently amended) The composition Composition according to claim 7, characterised in that wherein, said lecithin is present in said non-oily emulsion in an amount of 2-15 % (w/v), said bile salt is present in said non-oily emulsion in an amount of 2-15 % (w/v), and said cholesterol is

present in said non-oily emulsion in an amount of 2-15 % (w/v), in relation to the non-oily emulsion.

9. (currently amended) The composition Composition according to claim 7, or 8, characterised in that wherein the ratio by weight of lecithins, bile salts and cholesterol is 2:1:1.
10. (currently amended) The composition Composition according to claim 8, wherein any one of the proceeding claims, characterised in that the sum of the amounts of lecithins, bile salts and cholesterol constitutes 6-30 % (w/v) of the non-oily emulsion.
11. (currently amended) The composition Composition according to any one of the proceeding claims, characterised in that claim 1, wherein the composition further contains comprises an organic sulfur compound.
12. (currently amended) The composition Composition according to claim 11, wherein, characterised in that the organic sulfur compound is present in said composition in an amount of 2-30 % (w/v) and preferably in an amount of 4-25 % (w/v), in relation to the non-oily emulsion.
13. (currently amended) The composition Composition according to claim 11, wherein or 12, characterised in that the organic sulfur compound is selected from the group comprising consisting of dimethylsulfoxide, methylsulfonylmethane, 2,3-dimethylsulfolane, 2,4-dimethylsulfolane and sodium lauryl sulfate.
14. (currently amended) Use of the composition according to any one of claims 1 to 13 claim 1 for the manufacture of a cream, gel, lotion, suppositories, ointment, patch (TTS) for transdermal administration of active substances, preferably nutrients and/or medications.
15. (currently amended) Use of the composition according to any one of claims 1 to 13 claim 1 for transdermal administration of active substances, preferably nutrients and/or medications.
16. (new) The composition according to claim 12, wherein the organic sulphur compound is present in said composition in an amount of 4-25% (w/v), in relation to said non-oily emulsion.